(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COGERATION TREATY (PCT)

AIPO OMPL /5/ 0896×7

10/089687

(19) World Intellectual Property Organization International Bureau

(43) International Publication Date 5 April 2001 (05.04.2001)

PCT

(10) International Publication Number WO 01/23558 A3

(51) International Patent Classification²: C12N 15/12, C07K 14/705, C12Q 1/68, G01N 33/50, A01K 67/027

(21) International Application Number: PCT/US00/25610

(22) International Filing Date:

19 September 2000 (19.09.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

60/156.624	28 September 1999 (28.09.1999)	US
60/156,625	28 September 1999 (28.09.1999)	US
60/168.611	2 December 1999 (02.12.1999)	US
60/168,613	2 December 1999 (02.12.1999)	US
60/168,614	2 December 1999 (02.12.1999)	US

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier applications:

US	60/156,624 (CIP)
Filed on	28 September 1999 (28.09.1999)
US	60/156,625 (CIP)
Filed on	28 September 1999 (28.09.1999)
US	60/168,614 (CIP)
Filed on	2 December 1999 (02.12.1999)
US	60/168.611 (CIP)
Filed on	2 December 1999 (02.12.1999)
US	60/168,613 (CIP)
Filed on	2 December 1999 (02.12.1999)

(71) Applicant (for all designated States except US): INCYTE GENOMICS, INC. [US/US]; 3160 Porter Drive. Palo Alto, CA 94304 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): HODGSON, David, M. [US/US]; 567 Addison Avenue, Palo Alto. CA 94304 (US). LINCOLN, Stephen, E. [US/US]; 725 Sapphire Street, Redwood City. CA 94061 (US). RUSSO, Frank, D. [US/US]; 1583 Courdillaeras Road, Redwood City. CA 94062 (US). SPIRO, Peter, A. [US/US]; 3875 Park Boulevard, Apt. B16, Palo Alto. CA 94306 (US). BANVILLE,

Steven, C. [US/US]: 604 San Diego Avenue, Sunnyvale, CA 94086 (US). BRATCHER, Shawn, R. [US/US]: 550 Ortega Avenue #B321, Mountain View, CA 94040 (US). DUFOUR, Gerard, E. [US/US]; 5327 Greenridge Road. Castro Valley, CA 94552-2619 (US). COHEN, Howard, J. [US/US]: 3272 Cowper Street, Palo Alto, CA 94306-3004 (US). ROSEN, Bruce, H. [US/US]; 177 Hanna Way, Menlo Park, CA 94025 (US). SHAH, Purvi [IN/US]; 859 Salt Lake Drive, San Jose, CA 95133 (US). CHALUP, Michael, S. [US/US]; 183 Acalanes Drive. Apt. 6. Sunnyvale. CA 94086 (US). HILLMAN, Jennifer, L. [US/US]: 230 Monroe Drive, #17. Mountain View, CA 94040 (US). JONES, Anissa, Lee [US/US]: 445 South 15th Street, San Jose, CA 95112 (US), YU, Jimmy, Y. [US/US]; 37330 Portico Terrace, Fremont, CA 94536-7901 (US). GREENAWALT, Lila, B. [US/US]; 1596 Ballantree Way, San Jose, CA 95118-2106 (US). PANZER, Scott, R. [US/US]: 965 East El Camino, #621, Sunnyvale, CA 94087 (US). ROSEBERRY, Ann, M. [US/US]; 725 Sapphire Street, Redwood City, CA 94061 (US). WRIGHT, Rachel, J. [NZ/US]; 339 Anna Way, Mountain View, CA 94043 (US). CHEN, Wensheng [CN/US]; 210 Easy Street, #25, Mountain View, CA 94043 (US). LIU, Tommy, F. [US/US]: 201 Ottilia Street, Daly City, Ca 94014 (US). YAP, Pierre, E. [US/US]; 201 Happy Hollow Court. Lafayette. CA 94549-6243 (US). STOCKDREHER, Theresa, K. [US/US]; 1596 Ontario Drive, #2, Sunnyvale, CA 94087 (US). AMSHEY, Stefan [US/US]; 1541 Canna Court, Mountain View, CA 94043 (US). FONG, Willy, T. [US/US]: 572 Cambridge Street, San Francisco, CA 94134 (US).

- (74) Agents: HAMLET-COX, Diana et al.; Incyte Genomics. Inc., 3160 Porter Drive, Palo Alto, CA 94304 (US).
- (81) Designated States (national): AE. AG, AL. AM. AT. AU. AZ. BA. BB, BG. BR. BY. BZ, CA. CH. CN. CR. CU. CZ. DE. DK, DM. DZ. EE. ES, FI, GB. GD. GE, GH. GM. HR. HU. ID. IL, IN, IS. JP, KE. KG. KP, KR. KZ. LC. LK. LR. LS. LT. LU, LV, MA, MD. MG. MK. MN. MW. MX. MZ. NO. NZ, PL. PT, RO. RU, SD, SE, SG, SI, SK, SL, TJ, TM. TR. TT. TZ. UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH. GM. KE, LS, MW. MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European

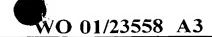
[Continued on next page]

(54) Title: HUMAN SECRETORY MOLECULES

(57) Abstract: The present invention provides purified secretory polynucleotides (sptm). Also encompassed are the polypeptides (SPTM) encoded by sptm. The invention also provides for the use of sptm, or complements, oligonucleotides, or fragments thereof in diagnostic assays. The invention further provides for vectors and host cells containing sptm for the expression of SPTM. The invention additionally provides for the use of isolated and purified SPTM to induce antibodies and to screen libraries of compounds and the use of anti-SPTM antibodies in diagnostic assays. Also provided are microarrays containing sptm and methods of use.

01/02/20

WO 01/23





patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

with international search report

(88) Date of publication of the international search report: 21 February 2002

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.





International Application No PCT/US 00/25610

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C12N15/12 C07K14/705 C12Q1/68 G01N33/50 A01K67/027 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) IPC 7 C12N C07K C12Q G01N A01K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Category ' Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. Х DATABASE EM_EST [Online] 1 - 19**EMBL** ID HSA29735, AC AA029735, 20 August 1996 (1996-08-20) HILLIER L ET AL.: "ze95c03.r1 Soares fetal heart NbHH19W Homo sapiens cDNA clone IMAGE: 366724 5', mRNA sequence." XP002162283 Note: 97.3 % nt sequence identity with SEQ ID NO:1 in 669 nt overlap. the whole document -/--Χ Further documents are listed in the continuation of box C. X Patent family members are listed in annex. Special categories of cited documents: T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance invention earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) involve an inventive step when the document is taken alone document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled in the art. other means document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 1 5.06. 01 8 March 2001 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016 van de Kamp, M

Form PCT/ISA/210 (second sheet) (July 1992)

3



International Application No
PCT/US 00/25610

Catalon of document, with indication, where appropriate, of the relevant passages Relevant to claim No.	C.(Continua	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	
EMBL ID AW009420, AC AW009420, 13 September 1999 (1999-09-13) STRAUSBERG R: "ws82d11.x1 NCI CGAP Co3 Homo sapiens cDNA clone IMAGE:2504469 3', mRNA sequence." XP002162284 Note: 99.6 % nt sequence identity with SEQ ID NO:1 in 547 nt overlap. the whole document A W0 99 03990 A (FLORENCE KIMBERLY A ;HUMAN GENOME SCIENCES INC (US); FENG PING (US) 28 January 1999 (1999-01-28) page 2, line 1-8 claims 1-23 A W0 99 18204 A (KATO SEISHI ;PROTEGENE INC (JP); SEKINE SHINGO (JP); SAGAMI CHEM R) 15 April 1999 (1999-04-15) page 3, line 1-17 claims 1-6 A W0 99 31236 A (BOUGUELERET LYDIE ;GENSET SA (FR); DUCLERT AYMERIC (FR); DUMAS MIL) 24 June 1999 (1999-06-24) page 3, line 23 -page 10, line 15	Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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(JP); SEKINE SHINGO (JP); SAGAMI CHEM R) 15 April 1999 (1999-04-15) page 3, line 1-17 claims 1-6 A WO 99 31236 A (BOUGUELERET LYDIE ;GENSET SA (FR); DUCLERT AYMERIC (FR); DUMAS MIL) 24 June 1999 (1999-06-24) page 3, line 23 -page 10, line 15	A	GENOME SCIENCES INC (US); FENG PING (US) 28 January 1999 (1999-01-28) page 2, line 1-8	1-19
SA (FR); DUCLERT AYMERIC (FR); DUMAS MIL) 24 June 1999 (1999-06-24) page 3, line 23 -page 10, line 15	A	(JP); SEKINE SHINGO (JP); SAGAMI CHEM R) 15 April 1999 (1999-04-15) page 3, line 1-17	1-19
	A	SA (FR); DUCLERT AYMERIC (FR); DUMAS MIL) 24 June 1999 (1999-06-24) page 3, line 23 -page 10, line 15	1-19

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

3





INTERNATIONAL SEARCH REPORT

International application No. PCT/US 00/25610

Box I O	bservations where certain claims were found unsearchable (Continuati n of item 1 of first sheet)
This Interna	ational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
	aims Nos.: ecause they relate to subject matter not required to be searched by this Authority, namely:
be	aims Nos.: scause they relate to parts of the International Application that do not comply with the prescribed requirements to such sextent that no meaningful International Search can be carried out, specifically:
	aims Nos.: ecause they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II O	bservations where unity of invention is lacking (Continuation of item 2 of first sheet)
1. A	is all required additional search fees were timely paid by the applicant, this International Search Report covers all
se	s all required additional search fees were timely paid by the applicant, this International Search Report covers all carchable claims.
2. As	all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment any additional fee.
3. As	s only some of the required additional search fees were timely paid by the applicant, this International Search Report overs only those claims for which fees were paid, specifically claims Nos.:
	o required additional search fees were timely paid by the applicant. Consequently, this International Search Report is stricted to the invention first mentioned in the claims; it is covered by claims Nos.: laims 1-19 (all partially)
Remark on	Pr test The additional search fees were accompanied by the applicant's protest. No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: Invention 1, claims 1-19 (all partially)

An isolated polynucleotide selected from the group consisting of a polynucleotide consisting of, or comprising the polynucleotide sequence of SEQ ID NO:1, homologs and variants, complementary polynucleotides, fragments and RNA equivalents of said polynucleotide. A composition for the detection of the expression of secretory polynucleotides comprising a polynucleotide as said, and methods of detecting said polynucleotide. A recombinant polynucleotide comprising a polynucleotide as said, operably linked to a promoter sequence, a cell and a transgenic organism comprising said recombinant polynucleotide, and a method for producing a secretory polypeptide by expressing said recombinant polynucleotide. A purified secretory polypeptide encoded by a polynucleotide comprising the polynucleotide sequence of SEQ ID NO:1, and an antibody binding to said polypeptide. A method of identifying a test compound which binds to said polypeptide. A microarray comprising a polynucleotide comprising at least 60 contiguous nucleotides of a polynucleotide as said, and a method for generating a transcript image using said microarray. A method for screening a compound for effectiveness in altering expression of a polynucleotide as said, and a method for assessing toxicity of a test compound using a probe comprising at least 20 contiguous nucleotide of a polynucleotide as said.

2. Claims: Inventions 2-63, claims 1-19 (all partially)

As invention 1, but concerning SEQ ID NOs 2-63, respectively.

International Application No PCT/US 00/25610

Information on patent family members

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